

MODIFIED ABSORPTION FEATURES OF TITANIA-ERBIUM  
INCORPORATED PLASMONIC TELLURITE GLASS SYSTEM

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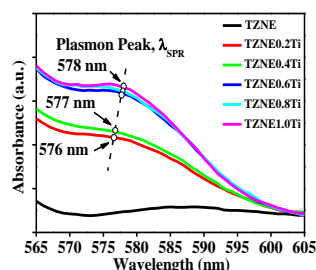
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## Graphical abstract



## Abstract

Achieving efficient lasing glass materials with enhanced absorption and emission cross-section by reducing the Rare Earth (RE) concentration quenching is a challenging issue. Metal nanoparticles (NPs) together with RE ions in the glass matrix are thought as a suitable alternative to overcome the limitations of concentration quenching and weak absorption of inorganic glasses. We prepare a series of Titania-Erbium doped Tellurite glass system with the form  $(69-x)\text{TeO}_2\text{-}20\text{ZnO-}10\text{Na}_2\text{O-}1\text{Er}_2\text{O}_3\text{-(x)TiO}_2$ , where  $0 \leq x \leq 1.0$  mol% via melt-quenching method with optimum erbium contents and varying  $\text{TiO}_2$  NPs concentrations. The NPs concentration dependent modifications in the absorption characteristics are scrutinized. Glasses are characterized via UV-Vis-NIR and XRD measurements. XRD pattern verifies the amorphous nature of prepared samples. The incorporation of  $\text{TiO}_2$  NPs is demonstrated to enhance the absorption intensity significantly. This augmentation is attributed to the effect of Surface Plasmon Resonance (SPR) mediated strong local electric field that is swallowed by neighboring  $\text{Er}^{3+}$  ions. The observed modification in optical energy band gap and Urbach energy are ascribed to the strong electric field around NPs that interact with the ligand of glass network to transform weak bond into defects. This observation is useful for the development of plasmonic nanoglass materials applicable for photonic devices.

Keywords: Tellurite glass, titanium dioxide nanoparticle, absorption spectra

## Abstrak

Untuk mencapai bahan kaca pelaseran yang efisien dengan penambahan daya penyerapan dan pancaran serta mengurangkan masalah penurunan keamatan pancaran Rare earth (RE) dalam kaca merupakan isu yang mencabar untuk ditangani. Logam Nanopartikel (NPs) bersama-sama dengan ion RE dalam matrik kaca dianggap alternatif yang sesuai untuk mengatasi masalah penyerapan RE yang rendah dan penurunan pendarkilau RE dalam kaca. Satu siri kaca Titania-Erbium yang didopkan dalam kaca Tellurite telah dihasilkan dengan komposisi  $(69-x)\text{Te}_2\text{O-}20\text{ZnO-}10\text{Na}_2\text{O-}1\text{Er}_2\text{O}_3\text{-(x)TiO}_2$ , dalam julat  $0 \leq x \leq 1.0$  mol% melalui kaedah pelindapkejukan-leburan dengan kandungan erbium yang optimum dan komposisi  $\text{TiO}_2$  NPs yang pelbagai. Kadar penyerapan bahan kaca mengandungi komposisi  $\text{TiO}_2$  NPs yang berbeza-beza diteliti. Kaca ini telah dicirikan menggunakan UV-Vis-NIR spektroskopi dan XRD. XRD telah mengesahkan sampel adalah bahan amorphous. Penambahan komposisi  $\text{TiO}_2$  NPs didapati mampu meningkatkan kadar penyerapan kaca. Peningkatan penyerapan ini adalah disebabkan Resonans Permukaan Plasmon (SPR) yang menghasilkan medan elektrik setempat dan memindahkan tenaga kepada RE. Jurang jalur tenaga optik dan tenaga Urbach diamati. Parameter ini dipengaruhi oleh medan elektrik di sekitar NPs dalam rangkaian kaca dimana ikatan yang lemah akan diubah menjadi kecacatan. Kajian ini dipercayai bermafaat bagi memperluaskan penggunaan bahan kaca plasmonik dan bahan peranti fotonik

Kata kunci: Kaca tellurite, titanium dioxide nanopartikel, spectra penyerapan

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## 1.0 INTRODUCTION

Nowadays, the demand of rare earth (RE) doped glasses is exponentially growing especially for photonic applications including cheap solid state laser, optical amplifiers, non-linear optical devices, and production of optical fibers. Selecting host glasses for RE ions are important for efficient up-conversion emission [1]. Tellurite based-glass is the most promising host glass due to several special feature such as wide transmission range from UV to IR region (340nm-15 $\mu$ m), low dispersion, superior transparency with refractive index between 2 to 2.5, high earth ion solubility [2], good thermal stability, chemical durability and low melting point [3].

Meanwhile, erbium remains the most attractive luminescent rare-earth element due to its emission in visible region which is beneficial for laser development [4]. However, Er<sup>3+</sup> has been disqualified to be nominated as strong emitter because of its small absorption and emission cross-section [5]. Several methods are proposed to overcome this drawback. The addition of second dopant either a REs or a transition metal which can enhance the optical properties of RE-doped glasses is suggested. Incorporation of metallic NPs into the glass are striking due to surface plasmon resonance (SPR) mediated impact on RE ions [6]. Apart from NPs manipulation and formation, glasses are the suitable dielectric host for activating enough RE ions via metallic NPs that act as sensitizers.

Silver and gold NPs is usually used as second dopant in Erbium-doped tellurite glass due to its chemical stability and strong SPR effect. However due to the wide range of parameters for separate applications, it is impossible to address all the requirements by use of silver and gold; therefore, alternative materials are needed to optimize the performance. Titanium NPs is attractive yet less exploited as second dopant in Erbium-doped tellurite glass even it exhibit SPR effect [7]. Metal oxide, TiO<sub>2</sub> NPs in the erbium-tellurite glass host is thought to offer high linear optical response [8] since the empty d-shell of Ti ion contribute strong linear polarizability [9]. Its oxide layer has advantage such as the position of SPR can be tuned by varying the thickness of oxide layer.

In addition, certain value of thickness of oxide layer can increase the SPR intensity instead of decreasing it. This effect make TiO<sub>2</sub> NPs more attractive for plasmonic and photonic application like solid-state laser [10]. Despite few systematic studies on titania-erbium doped tellurite glass, the mechanism of TiO<sub>2</sub> NPs stimulated spectroscopic modification is far from being understood.

We report the improvements in the absorption properties of TiO<sub>2</sub> NPs embedded erbium-doped tellurite glasses. TiO<sub>2</sub> NPs concentration dependent density, molar volume, optical energy band gap, Urbach energy, refractive index, and polarizability are determined. The synthesized disordered plasmonic media is demonstrated to be ideal for sundry nanophotonic applications.

**Table 1** Compositions (in mol%) of synthesized glass samples and their codes

Glass	TeO <sub>2</sub>	ZnO	Na <sub>2</sub> O	Er <sub>2</sub> O <sub>3</sub>	TiO <sub>2</sub>
TZNE	69	20	10	1	0
TZNE0.2Ti	68.8	20	10	1	0.2
TZNE0.4Ti	68.6	20	10	1	0.4
TZNE0.6Ti	68.4	20	10	1	0.6
TZNE0.8Ti	68.2	20	10	1	0.8
TZNE1.0Ti	68	20	10	1	1.0

## 2.0 EXPERIMENTAL

Series of glasses (enlisted in Table 1) with composition (69-x)TeO<sub>2</sub>-20ZnO-10Na<sub>2</sub>O-1Er<sub>2</sub>O<sub>3</sub>-(x)TiO<sub>2</sub> (where x ranges within 0 to 1.0 mol%) are synthesized using melt-quenching technique. Low doping level of Er<sup>3+</sup> is maintained to avoid undesirable attenuations such as dipolar interaction and luminescence quenching. Analytical grade starting materials of TeO<sub>2</sub> (purity 99.5%), ZnO (purity 99.9%), Na<sub>2</sub>O (purity 80.0%), Er<sub>2</sub>O<sub>3</sub> (purity 99.9%) and TiO<sub>2</sub> (purity 99.7%) from Sigma Adrich are acquired as glass constituents and thoroughly grinded. A platinum crucible holding this constituents mixture are placed inside an electrical furnace at

900°C for 20 minutes to achieve complete molten state. The melt is poured in a brass mould before transferring to an annealing furnace at 300 °C for 3 hours to remove residual thermal and mechanical strains. The sample is then cooled down to room temperature and stored in a desiccator to prevent a moisture attack. Finally, they are cut and polished to a thickness 2.5 $\pm$ 0.5 mm to produce shiny and scratch free surface for optical measurements.

The X-ray diffraction (XRD) measurements are performed using PANalytical X'Pert PRO MRD PW3040 with Cu Ka radiations ( $\lambda$  = 1.54 Å) in scanning angle of 2 $\theta$  ranging between 10° and 80° operated at 40kV and 35 mA. Room temperature optical absorption spectra

are recorded in the wavelength range of 200–1800 nm using a Shimadzu UVPC-3101 spectrophotometer with resolution of  $\pm 1$  nm. The glass density is measured by Archimedes method via Analytical balance of specific density-Precisa XT 220A with distilled water as an immersion liquid. The density  $\rho$  (g/cm<sup>3</sup>) of each sample is determined using the expression [11],

$$\rho = \rho_w \frac{W_a}{W_a - W_w} \quad (1)$$

where  $\rho_w$  is the density of distilled water (0.9975 g/cm<sup>3</sup>)  $W_a$  and  $W_w$  are the weight of the sample in the air and in water respectively.

The molar volume  $V_m$  is calculated using [12],

$$V_m = \sum_i \frac{x_i M_i}{\rho} \quad (2)$$

where  $x_i$  and  $M_i$  denote the molar fraction and molecular weight of the  $i$ th component, respectively. The refractive index  $n$  of glasses are calculated following [13],

$$\frac{n^2 - 1}{n^2 + 2} = 1 - \sqrt{\frac{E_{opt}}{20}} \quad (3)$$

where  $E_{opt}$  is the optical band gap energy of the glass obtained from the absorption data. The molar refractivity  $R_m$  is obtained using [14],

$$R_m = \frac{n^2 - 1}{n^2 + 2} (V_m) \quad (4)$$

The polarizability  $\alpha_e$  of the glasses is related to the refractive index through Lorentz-Lorenz equation and can be determined from [15],

$$R_m = \frac{4}{3} N_a \alpha_e \quad (5)$$

where  $N_a$  is the Avogadro's number.

The optical absorption coefficient  $\alpha(\lambda)$  is calculated from the following expression [16],

$$\alpha(\lambda) = \frac{2.303A}{d} \quad (6)$$

where  $A$  is absorbance and  $d$  is the thickness of the samples. The optical energy band gaps for direct and indirect transitions and the Urbach energy are calculated using the UV absorption edge. The frequency dependent absorption coefficient  $\alpha(\omega)$  yields [17],

$$\alpha(\omega) = \frac{B(\hbar\omega - E_{opt})^n}{\hbar\omega} \quad (7)$$

where  $B$  is a constant,  $E_{opt}$  is the optical band gap,  $\alpha(\omega)$  is the absorption coefficient at an angular frequency  $\omega = 2\pi\nu$ ,  $\hbar$  is the Planck constant divided by  $2\pi$  and  $n$  is the index can assume values of 0.5, 1.5, 2 and 3 depending on the nature of electronic transitions responsible for the absorption.

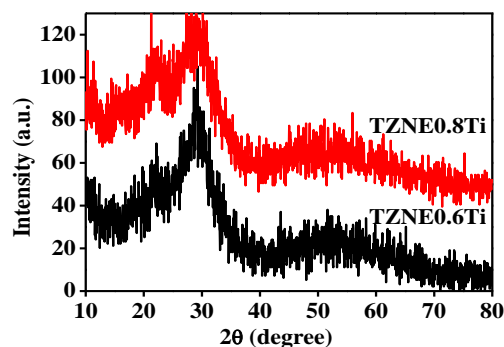
Urbach energy that characterizes the extent of the exponential tail of the absorption edge is given by [18]

$$\alpha(\omega) = B \exp\left(\frac{\hbar\omega}{\Delta E}\right) \quad (8)$$

where  $\Delta E$  is Urbach energy.

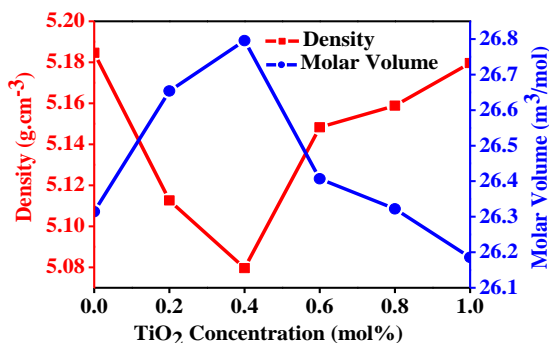
### 3.0 RESULTS AND DISCUSSION

Figure 1 displays the typical XRD spectra of two Er<sup>3+</sup> doped zinc-sodium tellurite glass samples containing titanium dioxide NPs. The presence of broad hump between 25° and 35° without any sharp crystallization peaks confirms the amorphous nature of these samples.



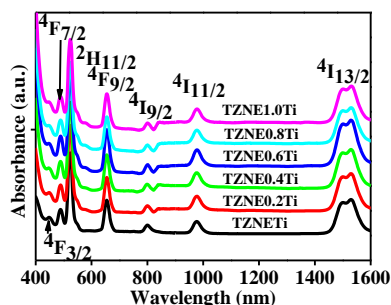
**Figure 1** Typical X-ray diffraction pattern for two samples with different TiO<sub>2</sub> NPs content

Figure 2 illustrates the TiO<sub>2</sub> NPs concentration dependent variation of density and molar volume. The density is found to decrease with the increase of TiO<sub>2</sub> NPs contents up to 0.4 mol% and then increased thereafter. This reduction in density is attributed to the substitution of higher molecular weight TeO<sub>2</sub> (159.60 g mol<sup>-1</sup>) compound by a lower molecular weight TiO<sub>2</sub> (79.876 g mol<sup>-1</sup>) entity in the glass network. The weak connectivity in the glass structure causes a drop in the density [19]. Furthermore, the increase in density is due to the increase of the glass network rigidity and compactness of the glass [15]. The increase in molar volume reflects the enhancement of free volume.

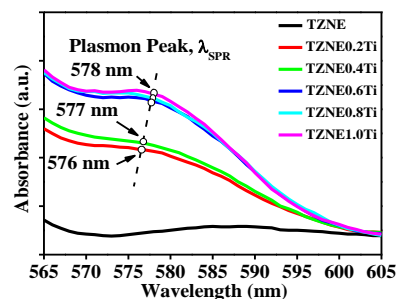


**Figure 2** TiO<sub>2</sub> NPs concentration dependent changes in density and molar volume

Figure 3 shows the UV-Vis-NIR absorption spectra of all samples. Seven absorption bands centered at 448, 489, 524, 655, 803, 978 and 1530 nm corresponding to the transition from the  $4I_{15/2}$  ground state to  $4F_{3/2}$ ,  $4F_{7/2}$ ,  $4H_{11/2}$ ,  $4F_{9/2}$ ,  $4I_{9/2}$ ,  $4I_{11/2}$  and  $4I_{13/2}$  excited state respectively are evidenced. The weak absorption band at 550 nm is assigned to  $4S_{3/2}$  transitions which overlap with upper neighboring level of  $4H_{11/2}$  [20]. Spectral features of all samples are qualitatively similar except the variation in intensity. Addition of TiO<sub>2</sub> NPs caused red shift at the absorption edge [24] which attributed to the formation of non-bridged oxygen (NBOs) [2]. In fact, a structural rearrangement of the glass network occurred with the increase of TiO<sub>2</sub> NPs concentration [25]. Even though the Er<sup>3+</sup> ions absorption bands overlapped the TiO<sub>2</sub> NPs resonance wavelength [26], however a weak plasmon band is detected in the range between 576–578 nm as shown in Figure 4. This weak band originated from TiO<sub>2</sub> NPs stimulated SPR effect [27]. Resonances of coherent oscillation of the surface conduction electron of TiO<sub>2</sub> NPs with the incident electromagnetic radiation manifested this plasmon absorption band [28]. Moreover, the SPR band can be tuned by varying the shape and size of NPs [29] which in usually done by accurately changing TiO<sub>2</sub> NPs concentration, annealing temperature and time.



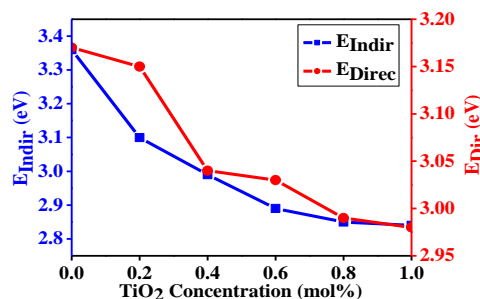
**Figure 3** UV-Vis-NIR absorption spectra of glass samples in the range of 400–1600 nm



**Figure 4** SPR band position of TiO<sub>2</sub> NPs

Optical transition and electronic band structure is analyzed using absorption spectra. Davis and Mott theory [17] is used to evaluate the optical band gap energy of the glass for both direct and indirect allowed transitions. In both transitions, the electromagnetic waves interact with the electrons in the valence band which are raised across the fundamental gap to reach the conduction band. However in amorphous material such as glass, indirect transition is most likely to occur and thus create band tail at the absorption edge. The Tauc's plot is drawn between  $(\alpha\hbar\omega)^n$  and photon energy ( $\hbar\omega$ ) by substituting  $n = 2$  for direct allowed transition and  $n = 1/2$  for indirect allowed transitions. The optical band gap values is obtained from the tangential intercept extrapolated at  $(\alpha\hbar\omega)^2 = 0$  and  $(\alpha\hbar\omega)^{1/2} = 0$  for direct and indirect transitions, respectively.

Figure 5 displays the TiO<sub>2</sub> NPs concentration dependent changes in the direct (3.17 – 2.98 eV) and indirect (3.36 – 2.84 eV). Reduction in optical band gap energy is due to the decrease in the average of bond energy and glass compactness. The addition of TiO<sub>2</sub> NPs in the host glass causes the formation of the NBO and reduces the BOs abundance. The decrease of optical band gap is due to low bonding energies of NBOs [30].



**Figure 5** Indirect and direct optical energy band gap with different TiO<sub>2</sub> NPs concentration

Figure 6 represents the Urbach energy as a function of NPs contents. The value of  $\alpha(\omega)$  near the absorption band edge exhibits an exponential behavior on the photon energy ( $\hbar\omega$ ) and obeys the empirical relation given by Urbach [16]. The exponential behavior of the band tail is due to the extensions of the valence and conduction bands to the band gap region. Figure 4(b) clearly shows increase in the band tail from 0.13 to 0.33 eV with the insertion of TiO<sub>2</sub> NPs into the glass. Urbach energy is used as yardstick to measure disordered nature of glass. Glass with large Urbach energy have greater tendency to convert weak bonds into defects due increase of disorder in the glass [31].

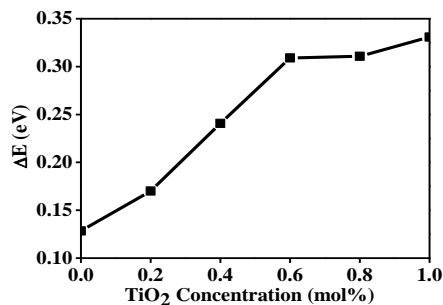


Figure 6 TiO<sub>2</sub> NPs concentration dependent Urbach energy

Figures 7 and 8 depict the TiO<sub>2</sub> NPs concentration dependent variation of refractive index and electronic polarizability, respectively. Introduction of TiO<sub>2</sub> increased the refractive index from 2.31 to 2.44. The increase of refractive index is interpreted in terms of the substitution of TeO<sub>4</sub><sup>4+</sup> ions with Ti<sup>4+</sup> ions. Addition of Ti<sup>4+</sup> ions caused the reconstruction of tellurite network and increased the NBOs atoms that have higher polarizability than BOs [30]. Therefore, incorporation of TiO<sub>2</sub> NPs at higher concentration has enhanced the refractive index of the glass. An overall increase in polarizability with the addition of Ti<sup>4+</sup> content is primarily ascribed to the creation of higher number of NBOs which have higher polarizability than BOs [8].

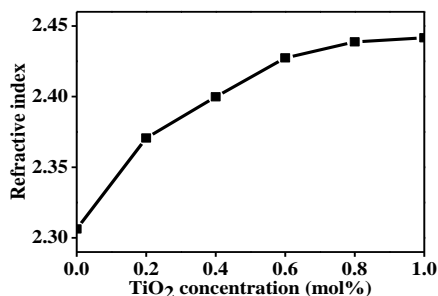


Figure 7 The refractive index against TiO<sub>2</sub> NPs concentration (mol %)

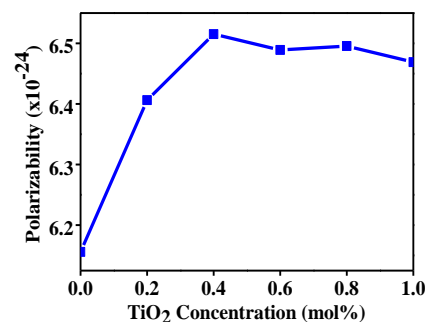


Figure 8 TiO<sub>2</sub> NPs concentration (mol %) dependent polarizability

## 4.0 CONCLUSION

TiO<sub>2</sub> NPs embedded erbium doped tellurite glasses acted as disordered plasmonic glass system are prepared and characterized. The impact of varying TiO<sub>2</sub> NPs concentration on physical and absorption properties are determined. The density and molar volume are found to vary in range of (5.18 - 5.07) g cm<sup>-3</sup> and (26.31 - 26.80) m<sup>3</sup> mol<sup>-1</sup>, respectively. SPR band of TiO<sub>2</sub> NPs is probed between 576 - 578 nm. The indirect and direct energy band gap are observed to decrease from (3.17 - 2.98) eV and (3.36 - 2.84) eV, respectively with the increase of TiO<sub>2</sub> NPs contents in the glass. The increase in Urbach energy in the range of (0.13 - 0.33) eV with the increase of TiO<sub>2</sub> NPs concentration is ascribed to the emergence of disorder via the creation of NBOs. Absorption features are observed to be modified with the incorporation of TiO<sub>2</sub> NPs. The improvement in absorption properties is attributed to the SPR mediated strong local field of TiO<sub>2</sub> NPs that appears in the close proximity of Er<sup>3+</sup> ions. Our study may provide useful information towards the development of photonic devices based on Titania-Erbium-Tellurite plasmonic disordered glass system.

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